

Citation for published version:

Wahila, MJ, Lebens-Higgins, ZW, Butler, K, Fritsch, D, Treharne, RE, Palgrave, RG, Woicik, JC, Morgan, B, Walsh, A & Piper, LFJ 2019, 'Accelerated optimization of transparent, amorphous zinc-tin-oxide thin films for optoelectronic applications', *APL Materials*, vol. 7, no. 2, 022509. <https://doi.org/10.1063/1.5053683>

DOI:

[10.1063/1.5053683](https://doi.org/10.1063/1.5053683)

Publication date:

2019

Document Version

Peer reviewed version

[Link to publication](#)

Publisher Rights

CC BY

University of Bath

Alternative formats

If you require this document in an alternative format, please contact:
openaccess@bath.ac.uk

General rights

Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

Take down policy

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Accelerated optimization of transparent, amorphous zinc-tin-oxide thin films for optoelectronic applications

Matthew J. Wahila,¹ Zachary W. Lebens-Higgins,¹ Keith T. Butler,² Daniel Fritsch,³ Robert E. Treharne,⁴ Robert G. Palgrave,⁵ Joseph C. Woicik,⁶ Benjamin J. Morgan,² Aron Walsh,⁷ and Louis F. J. Piper⁸

¹*Department of Physics, Applied Physics and Astronomy, Binghamton University, Binghamton, New York 13902, USA*

²*Department of Chemistry, University of Bath, Claverton Down, Bath BA2 7AY, United Kingdom*

³*Department Structure and Dynamics of Energy Materials, Helmholtz-Zentrum Berlin für Materialien und Energie, Hahn-Meitner-Platz 1, 14109 Berlin, Germany*

⁴*Stephenson Institute for Renewable Energy, University of Liverpool, Liverpool, L69 7ZF, United Kingdom*

⁵*University College London, Department of Chemistry, 20 Gordon Street, London, WC1H 0AJ, United Kingdom*

⁶*Materials Science and Engineering Laboratory, National Institute of Standards and Technology, Gaithersburg, Maryland 20899, United States*

⁷*Department of Materials, Imperial College London, Exhibition Road, London SW7 2AZ, United Kingdom*

⁸*Materials Science & Engineering, Binghamton University, Binghamton, New York 13902, USA^{a)}*

In the last decade, transparent amorphous oxide semiconductors (TAOS) have become an essential component of many electronics, from ultra high resolution displays to solar cells. However, these disordered oxides typically rely on expensive component metals like indium to provide sufficient charge carrier conduction, and their optoelectronic properties are not as predictable and well-described as those of traditional, crystalline semiconductors. Herein we report on our comprehensive study of the amorphous zinc-tin-oxide (a-ZTO) system for use as an indium-free, n-type TAOS. Using a combination of high-throughput co-deposition growth, high resolution spectral mapping, and atomistic calculations, we explain the development of disorder-related subgap states in SnO₂-like a-ZTO and optical bandgap reduction in ZnO-like a-ZTO. In addition, we report on a composition-induced electronic and structural transition in ZnO-like a-ZTO resulting in an exceptionally high Figure of Merit, comparable to that of amorphous indium-gallium-zinc-oxide (a-IGZO). Our results accelerate the development of a-ZTO and similar systems as indium-free TAOS materials.

Transparent amorphous oxide semiconductors (TAOS) are currently of great interest due to the wide range of optoelectronic applications enabled by their high transparencies and charge carrier mobilities combined with their potential for low temperature fabrication on inexpensive substrates.¹ TAOS systems that incorporate multiple cation species are particularly promising as they offer the ability to fine tune optoelectronic properties for specific applications by varying the atomic composition and coordination.^{2–7} Unfortunately, most TAOS systems investigated thus far incorporate rare, expensive, and/or toxic cations such as indium, hindering their development and reducing their practicality.⁸ Finding a material system with similar performance to the prototypical TAOS, amorphous indium-gallium-zinc-oxide (a-IGZO),² but composed of only inexpensive, abundant, and non-toxic elements would represent a substantial advancement in the development of TAOS materials.

Amorphous zinc-tin-oxide (a-ZTO), which possesses many desirable properties for thin film transistor applications, has recently arisen as a potential a-IGZO alternative.^{9,10} ZTO films contain only inexpensive and innocuous elements, and can be fabricated using a wide variety of industrially relevant methods, including sol-gel,¹¹ magnetron sputtering,^{7,12–15} chemical vapor,¹⁶ pulsed laser,¹⁷ and atomic layer depositions.^{9,18,19} Some researchers have even employed co-deposition techniques to quickly fabricate large sample sets and accelerate investigation of the large compositional phase space.^{7,12,20,21} However, most of these studies have focused on the optimization of films for specific device applications rather than attempting to explain the physical origins of the evolving material properties.

Recent research has focused predominantly on the effects of post growth anneal treatments, compositional variations, and material interactions on the properties of complete electronic devices, rather than investigating the underlying physics of the amorphous ZTO system.^{6,7,22–27} Several recent studies have begun to investigate the effects of composition and disorder on the electronic structure of ZTO;^{17,19,28} however, the full range of possible cation ratios has remained unexplored and the atomistic origin of many optical and electronic properties of these ZTO films remains uncertain.

^{a)}Electronic mail: lpiper@binghamton.edu

Herein, we present our study of the a-ZTO system using co-deposition of thin films in tandem with multiple optical and x-ray techniques, including ellipsometry and synchrotron-based hard x-ray photoelectron spectroscopy. Our high-resolution electronic and optical property mapping elucidates the evolution of electronic structure across a wide range of compositions, helping to identify and correlate key phenomena occurring across the a-ZTO compositional phase space. Combining our experimental results with density functional theory calculations, we then explain these phenomena in terms of the local structure and atomic coordination within these structurally disordered films.

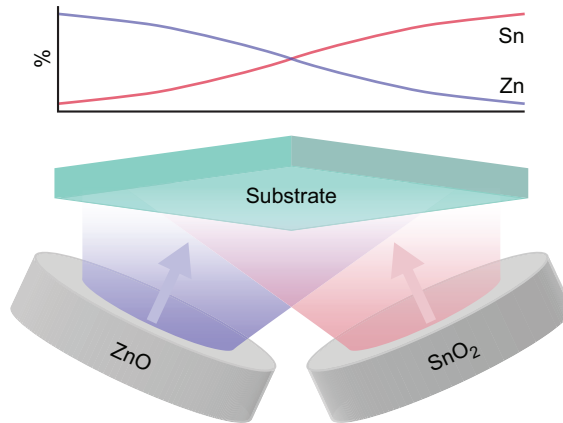


FIG. 1. **Co-deposition method.** Schematic showing how the co-deposition process creates a continuous compositional gradient across the film deposition area.

In order to thoroughly investigate the a-ZTO system, thin films were deposited on 10×10 cm OptiWhite soda-line glass by room temperature, magnetron sputtering co-deposition at University of Liverpool, as depicted in Figure 1. Powers of 250 and 80 W were applied to the ZnO and SnO₂ sputter guns, respectively, under a 5 mTorr Ar atmosphere. These large samples were then cut into smaller pieces for easier experimental characterization (see Supplementary Figure 1). This fabrication method produces a continuous compositional gradient across the deposition area, allowing for the simultaneous investigation of a large swath of the compositional phase space.^{7,12,20,21} Amorphous SnO₂ and polycrystalline ZnO films were fabricated for reference using single target depositions with the same parameters.

Figures 2 (a) - 2 (b) show the variation in cation ratio and film thickness across the 10×10 cm film resulting from the co-deposition. The compositional map, shown in Figure 2 (a), was created from scanning x-ray photoelectron spectroscopy (XPS) core-level spectra performed on 256 points across the sample area using a laboratory-based monochromated Al K α source with a hemispherical analyzer located at University College London. Measurements were performed in constant analyzer energy mode with a pass energy of 151.2 eV. The Sn:Zn ratio was found to vary smoothly from a minimum of 9.23% tin on the bottom-left to a maximum of 75.30% tin on the top-right.

The film thickness map, shown in Figure 2 (b), was created using ellipsometry measurements taken on an 81 point measurement grid across the central 8×8 cm² portion of the sample area using a J.A. Woollam Co. spectroscopic ellipsometer at University of Liverpool. The observed compositional gradient and thickness variations from this co-deposition compare well with trends predicted using ellipsometry on single target deposition films (see Supplementary Figures 2 and 3).

The optical band gap evolution, shown in Figure 2 (c), was determined using Tauc analysis of ultraviolet-visible spectroscopy (UV-Vis) measurements taken at 83 points across the sample area using either a Filmetrics F20 series reflectometer with T-1-UV transmittance upgrade or a Shimadzu UV-1800 UV-Vis spectrophotometer, both located at Binghamton University.²⁹ Additional analysis was performed on the UV-Vis spectra to estimate film thicknesses and indices of refraction for comparison with the ellipsometry results (see Supplementary Figure 4).^{30,31} Interestingly, the optical band gap evolution does not directly correspond to film composition, with the high tin percentage corner of the sample displaying an almost constant optical band gap independent of composition, while the low tin percentage corner displays a steep drop in band gap size with decreasing tin content (see Supplementary Figures 5).

Film resistivity, shown in Figure 2 (d), was calculated for 131 points across the sample area using sheet resistances measured with a 4-point probe system located in the Nanofabrication Facility at Binghamton University and film thicknesses calculated from ellipsometry (see Supplementary Figure 6). There are significant variations in resistivity across the sample which can be correlated to both the film thickness and Sn:Zn ratio. While film thickness does have an effect on the resistivity, it follows the expected trend of thinner regions displaying higher resistivity than thicker regions when Sn:Zn ratio is constant. The more interesting trends are attributable to the films' Sn:Zn ratio. While the high tin percentage corner of the sample has a fairly uniform resistivity between 10 - 100 Ω -cm, both the highest

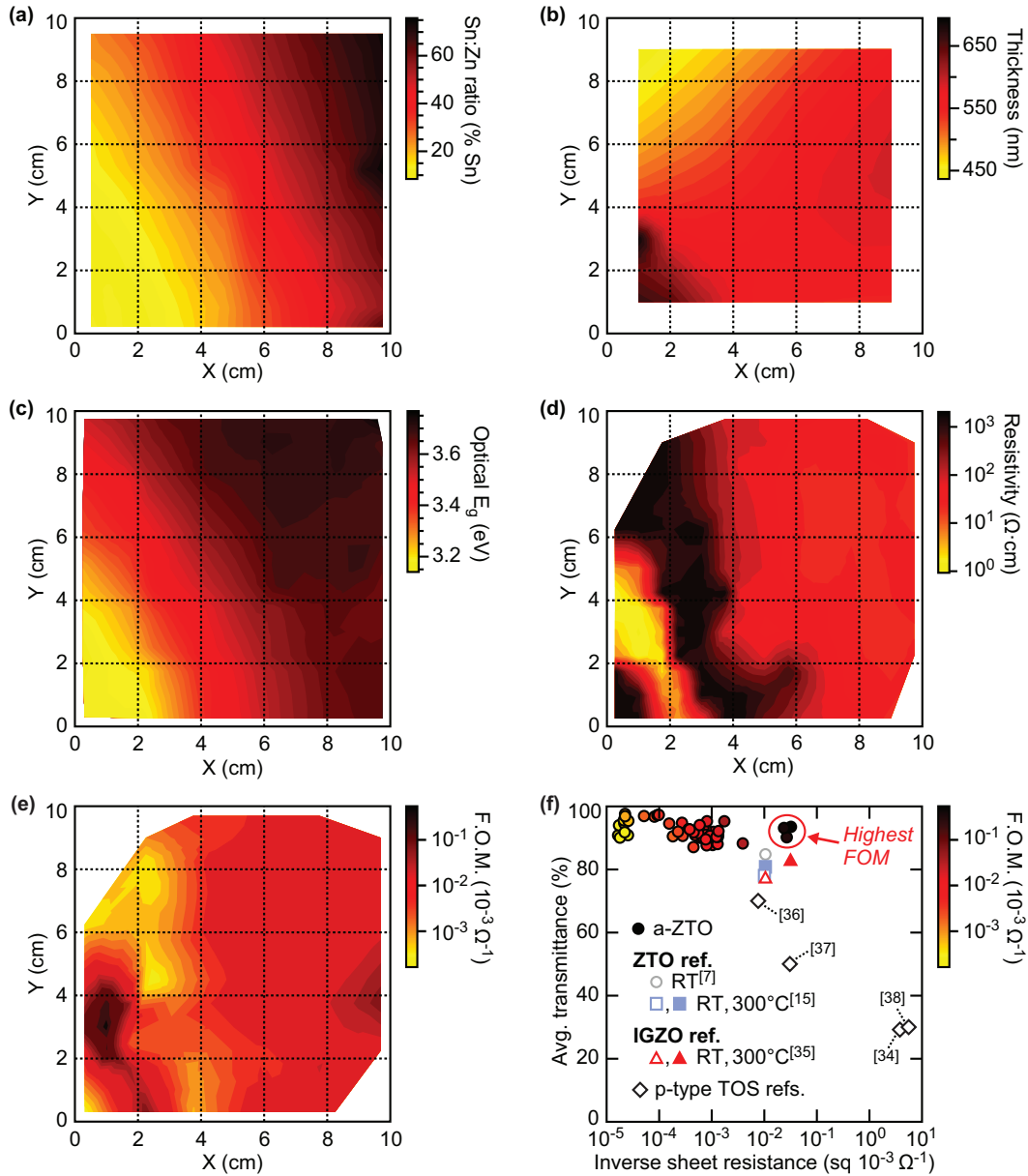


FIG. 2. **Film characterization across the co-deposition area.** Contour maps of **a**, Sn:Zn ratio, **b**, film thickness, **c**, direct optical band gap, **d**, resistivity, and **e**, Figure of Merit across the 10×10 cm a-ZTO thin film deposition area. **f**, Graphical representation of electrical resistance, optical transmission, and FOM for points across the a-ZTO film, including reference points from the literature for comparable ZTO (gray circle, blue squares) and IGZO (red triangles) sputtered thin films, as well as high-quality, crystalline p-type TAOS films (black diamonds).

(> 2000 $\Omega \cdot \text{cm}$) and lowest (< 1 $\Omega \cdot \text{cm}$) resistivities are found near the low tin percentage corner. Of particular interest is the region of ~ 10 - 12% tin, where film resistivity is drastically lower than neighboring regions with slightly more or less tin content.

Figure of Merit (FOM) values were calculated for points across the a-ZTO film, shown in Figure 2 (e). The FOM of a thin film can be calculated using the formula $\text{FOM} = \sigma / \alpha$, where σ is the film conductivity and α is the optical absorption coefficient. If optical reflectance is negligible, this can be approximated by $\text{FOM} \approx -1/R_S \ln(T)$, where R_S is the sheet resistance and T is the wavelength-averaged optical transmittance.³²⁻³⁴ Although optical transparency across the entire a-ZTO film is extremely high, much of the film is too resistive to result in a reasonable FOM. However, the previously mentioned region of ~ 10 - 12% tin possesses a low enough resistance to give FOM values comparable to or better than previously reported values for a-ZTO and other high FOM TAOS materials.^{7,15,35}

Shown in Figure 2 (f), the best resistivity values reported in the literature for a-ZTO have remained fairly constant

for many years.^{7,15} Surprisingly, our low tin percentage a-ZTO possess both better transparency and better resistivity than previously reported for high tin percentage a-ZTO films. Additionally, although Hall measurements are not well suited to investigate non-uniform and amorphous films, some Hall measurements were performed on select 1×1 cm film pieces. The results indicate that our low resistivity, low tin percentage region possesses comparable Hall mobility and bulk concentration to our high tin percentage region despite having a several orders of magnitude higher conductivity (see Supplementary Figure 7).

While higher temperature fabrication up to 300°C has been found to produce slightly increased transparency in high tin percentage a-ZTO films, it has not led to significantly improved resistivity.¹⁵ This puts high tin percentage a-ZTO at a distinct disadvantage to a-IGZO, which has been shown to have both improved electronic and optical properties following low-temperature processing.⁵ In contrast, our low tin percentage a-ZTO is found to already possess a better FOM than some a-IGZO reported in the literature which was annealed up to 300°C .³⁵

Additionally, our a-ZTO also compares favorably to transparent, crystalline p-type oxides such as CuAlO_2 ,³⁶ ZnRh_2O_4 ,³⁷ CuCrO ,³⁸ and SrCrO .³⁴ These crystalline p-type materials are able to match or even surpass amorphous n-types in terms of resistivity, but at the cost of significantly reduced optical transparency due to subgap states or disruption of the mechanisms responsible for their large optical band gaps. In contrast, while our low tin-percentage a-ZTO does possess a slightly smaller optical band gap than our high tin percentage film, the gap remains larger than 3 eV across the entire deposition area resulting in excellent optical transparency at all compositions.

From these results, we note two compositional regions warranting further investigation; the top-right corner possessing 60% tin or more (tin-rich), and the bottom-left corner possessing 30% tin or less (tin-poor). The tin-rich film possesses a large optical band gap and reasonable resistivity which both remain fairly constant over a large compositional range, indicating tin-rich a-ZTO could make a good TAOS material as suggested by other studies.^{7,21} In contrast, the tin-poor film possesses a rapidly changing optical band gap and both the highest and lowest observed resistivity. Understanding the origin of the anomalously low resistivity in this region could lead to the development of inexpensive ZTO-based materials for transparent conducting oxide (TCO) applications.^{39,40}

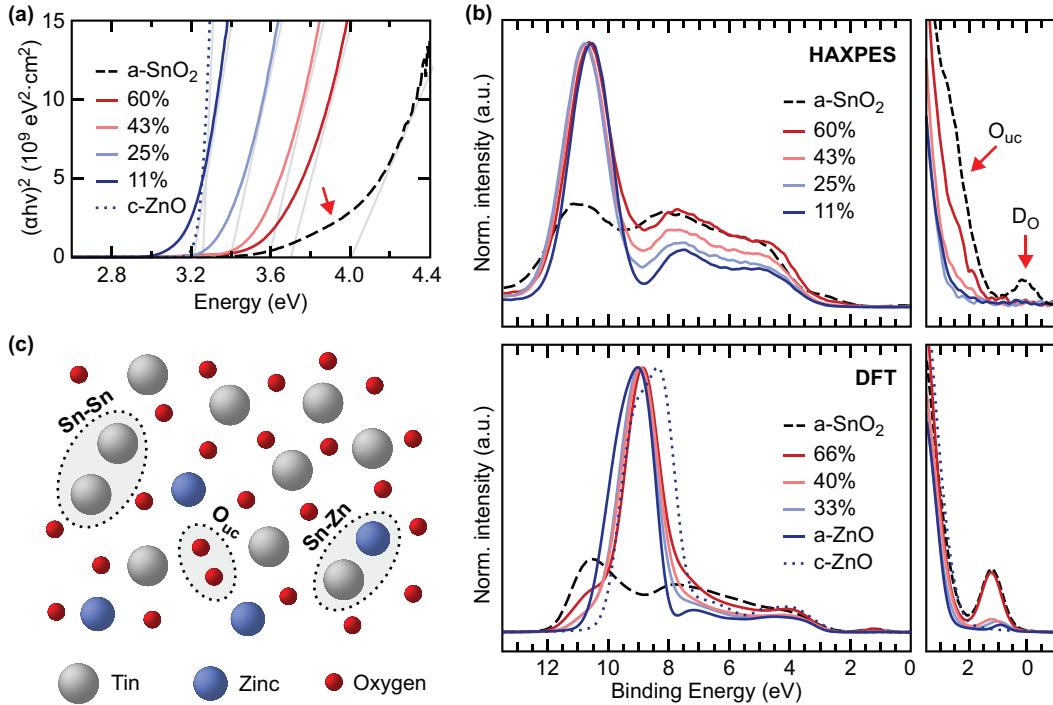


FIG. 3. Subgap state formation in tin-rich a-ZTO. **a**, Tauc plots for select points from high to low tin composition, including c-ZnO and a-SnO₂ for reference. **b**, HAXPES valence band spectra and DFT total density of states after weighting and convolution for several compositions of a-ZTO, including a-SnO₂ and c/a-ZnO for reference. Under-coordinated oxygen (O_{uc}) and oxygen deficiency (D_O) related subgap states are identified. **c**, Simplistic diagram of potential under-coordinated oxygen and oxygen deficiency defects within the a-ZTO.

Tauc plots for points along a diagonal between the tin-rich and tin-poor corners of the deposition area are shown in Figure 3 (a), including plots for reference amorphous SnO₂ (a-SnO₂) and polycrystalline ZnO (c-ZnO). For comparison, tauc plots were also created using optical constants calculated with hybrid DFT (see Supplementary Figure 8).⁴¹ The band gap of the a-ZTO film is found to increase as tin concentration increases, with the a-SnO₂ reference clearly

possessing the largest optical band gap. Additionally, a broadening of the absorption edge is observed in all of the a-ZTO and a-SnO₂ spectra, which is mostly consistent with exponential Urbach tail absorption due to structural disorder.^{42,43} However, the more tin-rich a-ZTO and a-SnO₂ display additional absorption below the main absorption edge (indicated by a red arrow), which is likely due to more than just Urbach tailing.

To investigate the tendency of tin-rich a-ZTO to form defect associated subgap states, hard x-ray photoelectron spectroscopy (HAXPES) measurements were performed at the National Institute of Standards and Technology (NIST) bending magnet beamline X24 at the National Synchrotron Light Source (NSLS) at Brookhaven National Laboratory. Measurements were performed at a ~ 4 keV photon energy, with a pass energy of 500 eV and a Gaussian instrumental broadening of 0.45 eV. Energy calibration of the HAXPES spectra was performed using external Au references in electrical contact with the samples.

For comparison, density functional theory calculations were performed for several stoichiometric a-ZTO compositions. Structures were first generated using a melt and quench approach, previously demonstrated to provide good agreement with experimental results.^{44,45} Starting from an initial crystal structure of each composition the system was heated to 3000 K using molecular dynamics (MD) and allowed to evolve for 10 ps, using a time step of 0.5 fs. Following the melting, the resultant systems were cooled to 0 K at a rate of 5×10^{13} K s⁻¹. The structures were then allowed to relax until the forces on all atoms were below a threshold value of 0.005 eV Å⁻¹.

The DFT calculations were performed in the Vienna Ab Initio Simulation Package (VASP)⁴⁶ within the projector augmented wave formalism,⁴⁷ using Gamma point sampling, the PBEsol functional,⁴⁸ and a cutoff energy of 400 eV. The final valence band density of states (DOS) were then produced by weighting each partial DOS with the appropriate photoionization cross-section, summing the partial DOS together, and then applying a pseudo-Voigt profile convolution to match experimental broadening. The calculated spectra were then energetically aligned to the experimental spectra for comparison. It must be noted that while the Zn 3d semi-core level peak position is clearly underestimated by the DFT, this is to be expected due to the lack of explicit on-site corrections in the calculations.⁴⁹ This misplacement of the Zn 3d should not significantly affect the calculated density of states near the valence and conduction band edges.

Both the HAXPES and DFT total DOS of the valence band, shown in Figure 3 (b), confirm the tendency of tin-rich a-ZTO to form defect associated subgap states. Both midgap states and near-edge states immediately above the valence band edge are observed for only the more tin-rich a-ZTO compositions, with HAXPES spectra for the tin-poor compositions showing no evidence of subgap states. These coordination-related subgap states are likely the cause of the additional optical absorption observed below the main absorption edge of the tin-rich compositions.

Previous theoretical and experimental work has confirmed that these subgap states originate from the local coordination of cation (tin/zinc) and anion (oxygen) atoms within the disordered structure, as depicted in Figure 3 (c).^{44,50-53} These defect states can exist in perfectly stoichiometric films provided there is enough variation in atomic coordination environments. Moreover, work on a-IGZO has shown that it is possible to modify or even remove these defect states by adjusting local atomic coordinations with only low temperature processing.⁵ The possibility of achieving a similar control of subgap states within tin-rich a-ZTO makes it an excellent candidate for further materials optimization.

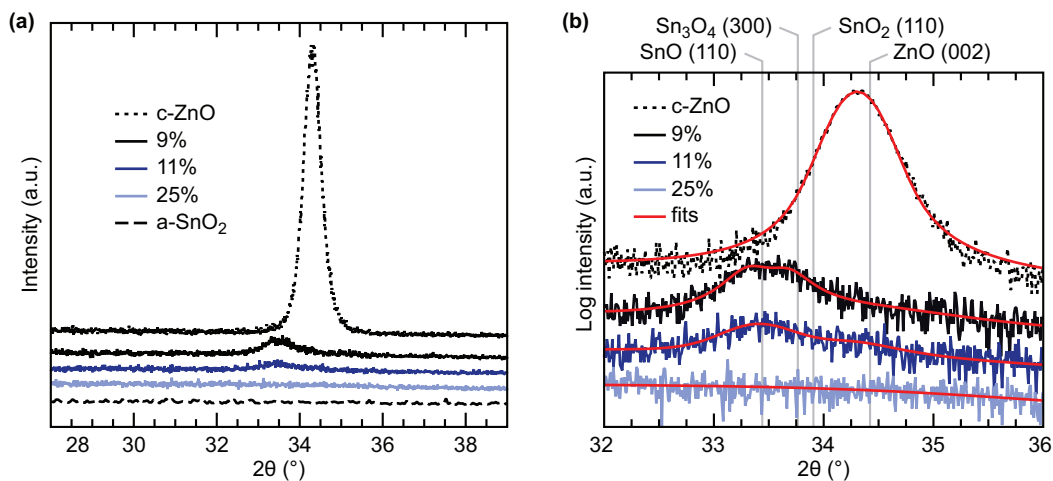


FIG. 4. **Crystallite formation in tin-poor a-ZTO.** **a**, XRD spectra for select points across the tin-poor corner, including a c-ZnO and a-SnO₂ references. Spectra have been truncated to only show the region containing observed features. **b**, Truncated XRD including curve fits.

On the other end of the compositional phase space, tin-poor a-ZTO displays intriguing behavior which has yet to be explored. In general, the tin-rich film possesses reasonable resistivity while the tin-poor film possesses very high resistivity, as shown in Figure 2 (d). However, the sudden drop in resistivity near 10 - 12% tin is an anomaly. This low resistivity gives the tin-poor a-ZTO film an unusually high FOM for an amorphous oxide semiconductor.³²⁻³⁴ One indication of why this compositional region displays unexpectedly low resistivity is given in Figure 4.

Figure 4 (a) shows x-ray diffraction (XRD) spectra taken on points across the tin-poor corner of the deposition area using a PANalytical X'Pert PRO XRD system located in the Analytical and Diagnostics Laboratory (ADL) at Binghamton University. A weak XRD feature near 33.5° is found to appear between ~ 25 - 11% tin, and grow more intense as tin percentage is further reduced. The c-ZnO reference film grown using the same deposition parameters exhibits a large ZnO (002) peak at a slightly higher diffraction angle, suggesting the feature observed in the tin-poor a-ZTO could be due to the formation of some strained ZnO crystallites.^{12,20} However, as shown in Figure 4 (b), this weak feature appears to be made up of more than one peak for at least some compositions and could also be due to segregated Sn_xO_y crystallites of different compositions within the tin-poor a-ZTO matrix.⁴⁴

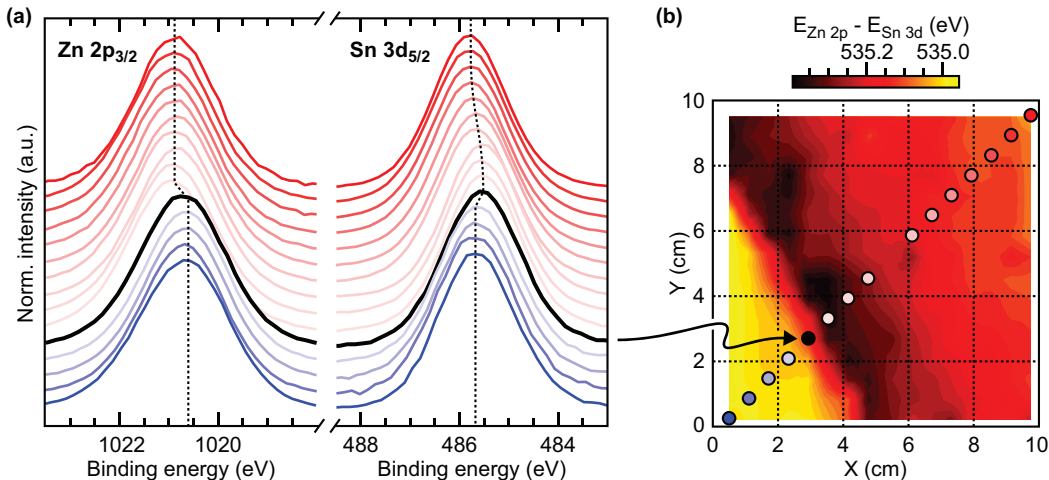


FIG. 5. **Peak shifts in tin-poor a-ZTO.** **a**, Example Zn 2p_{3/2} and Sn 3d_{5/2} XPS spectra for select points from high to low tin percentages. **b**, Contour map of the Zn 2p - Sn 3d core-level energetic separation across the entire deposition area.

Shown in Figure 5 (a), Zn 2p and Sn 3d XPS core levels taken from select spots across the deposition area display compositionally-dependent shifts that correlate with the observed crystallite formation. Commensurate shifts are also observed in the near-valence Zn 3d and Sn 4d levels as well as auger peaks measured using both high-resolution XPS and HAXPES (see Supplementary Figures 9, 10, and 11). The Zn 2p peak displays an abrupt energetic shift towards the Fermi level within the composition range where weak XRD peaks begin to appear. This shift indicates a sudden and sustained change in the local zinc bonding environment with the onset of crystallization, and agrees with the expected behavior of the Zn 3d upon crystallization as predicted by DFT, as shown in Figure 3 (b). Concurrently, the Sn 3d peak displays a gradual energetic shift towards the Fermi level as the film composition approaches the onset of crystallization, abruptly returning to its original energetic position once some weak crystallization is observed.

Figure 5 (b) maps the evolution of the Zn 2p and Sn 3d core level separation across the entire deposition area. Plotting core level separation rather than individual peak positions removes any contribution from Fermi level shifts or other confounding effects, providing clearer insight into changes of the local environment around cation species. The core level separation is found to gradually increase by ~ 0.3 eV between 40 to 20% tin, and then abruptly decrease in the ~ 25 - 11% tin range where initial crystallization is observed. This smaller separation is then maintained for the entire tin-poor corner of the film.

Figure 6 shows three important optoelectronic properties as a function of film Sn:Zn ratio, i.e., optical band gap, film resistivity, and energetic separation of key cation core levels. The evolution of these properties highlights important local disorder-related phenomena that must be taken into account when developing a-ZTO thin films for different applications.

The optical band gap evolution of disordered ZTO, shown in Figure 6 (a), displays band gap reduction at both compositional ends (indicated by the red arrows). The observed band gap changes are confirmed using both optical transmittance and ellipsometry measurements (see Supplementary Figure 12). Local coordination defects, which can play a large role in reducing optical transparency through subgap state absorption, are far more likely at SnO_2 -like (tin-rich) compositions due to the higher oxygen to cation ratio. These defect states and their corresponding optical absorption are clearly observed with increasing magnitude as tin content increases, up to and including the pure

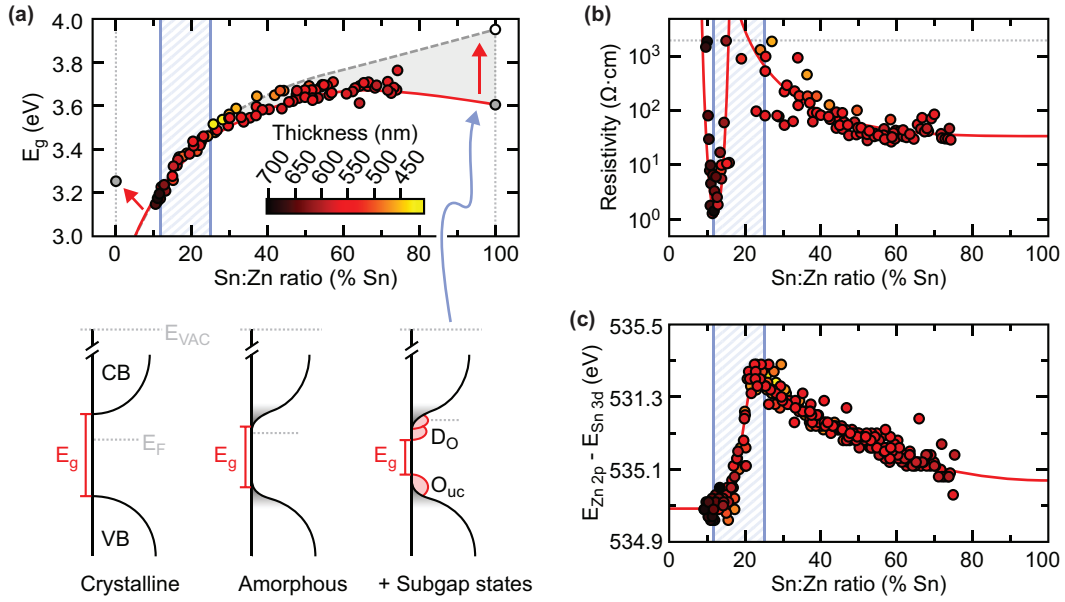


FIG. 6. **Summary of observed optoelectronic phenomena.** **a**, Evolution of the direct optical band gap with Sn:Zn ratio, including a-SnO₂ and c-ZnO endpoints. Band gap trends including (red line) and excluding (gray dashed line) subgap absorption are extrapolated to the a-SnO₂ endpoint determined by both including (gray circle) and excluding (white circle) weak absorption tailing. **b**, Evolution of film resistivity with Sn:Zn ratio, only including points with resistivity below the maximum value measurable by our equipment (dotted gray line). **c**, Evolution of the Zn 2p - Sn 3d energetic separation with Sn:Zn ratio. For all plots, trends are extrapolated across the full Sn:Zn ratio range (red line) and the compositional region wherein weak crystallization begins is indicated (blue hatching).

a-SnO₂ endpoint. The ability to control these defects is integral to the further development of tin-rich a-ZTO for applications where high optical transparency is desired.

While disordered ZnO-like (tin-poor) compositions can also suffer from band gap reduction compared to the c-ZnO endpoint, this is not attributed to under-coordinated oxygen- or cation-related defect states as it is for SnO₂-like compositions. Instead, first-principles simulations suggest the cause to be band edge changes resulting from altered Zn 4s - O 2p hybridization in the disordered phase (see Supplementary Figure 13). Disorder in pure ZnO has been suggested to decrease Zn 4s - O 2p hybridization, resulting in a lower intersite hopping probability and more localized Zn 4s conduction states.^{54,55} This localization would curtail percolation/hopping conduction through the film, greatly increasing resistivity.^{2,4} This is in contrast to a-SnO₂, where high mobilities comparable to that of the crystalline phase are known to be achievable.⁵⁶ Thus, more ZnO-like compositions might be expected to possess lower electron mobilities than more SnO₂-like compositions.

Figure 6 (b) shows the resistivity changes with composition, mostly confirming the expected trend of disordered SnO₂-like (tin-rich) compositions being far less resistive than the disordered ZnO-like (tin-poor) compositions. However, it also reveals that the anomalous drop in resistivity at certain low tin concentrations, which correlates with the observed onset of weak crystallization. Furthermore, this onset of weak crystallization strongly correlates with the sudden shifts in relative core-level peak positions measured using XPS/HAXPES, as shown in Figure 6 (c). Taken together, all of this data points towards a transition between two discrete material regimes with markedly different local atomic interactions occurring around a 20% tin to zinc ratio.

In summary, using thin film co-deposition in conjunction with high-resolution spatial mapping of film properties, we have correlated the observed evolution of optoelectronic properties with changes in the local electronic and atomic structures. In addition to clarifying the cause and effects of subgap state formation on film properties across a wide compositional range, we have also revealed a thus far unreported composition-induced electronic transition in tin-poor amorphous zinc-tin-oxide, which results in high Figure of Merit values comparable to room temperature grown a-IGZO.

While disorder in these multi-cation oxides typically results in poor charge carrier conduction, the observed electronic transition represents an unexpected exception. Contrary to conventional wisdom, whereby one might expect the greatest conduction to occur at tin-rich compositions due to the spherical tin 5s orbital contributions in the conduction band, the observed electronic and structural changes instead result in a narrow range of tin-poor compositions possessing a drastically increased conduction. Understanding these material behaviors, both expected and

unexpected, is critically important for the further development of amorphous zinc-oxide based semiconductors for electronics applications.

See supplementary material for additional details on the film growth method, details on the optical analysis, comparison between UV-Vis and ellipsometry results, comparison between theoretical and experimental Tauc plots, additional XPS/HAXPES spectra, and valence band partial density of states from DFT calculations.

Use of the National Synchrotron Light Source at Brookhaven National Laboratory, was supported by the U.S. Department of Energy, Office of Science, Office of Basic Energy Sciences, under Contract No. DE-AC02-98CH10886. This material is based upon work supported by the Air Force Office of Scientific Research under award number FA9550-18-1-0024. This research has received funding from the European Union's Horizon 2020 research and innovation program under grant agreement No. 641864 (INREP). This work made use of the ARCHER UK National Supercomputing Service (<http://www.archer.ac.uk>) via the membership of the UK's HPC Materials Chemistry Consortium, funded by EPSRC (EP/L000202) and the Balena HPC facility of the University of Bath. BJM acknowledges support from the Royal Society (UF130329). ZL-H was supported by a scholarship from the Summer Scholars and Artists Program at Binghamton University.

- ¹X. Yu, T. J. Marks, and A. Facchetti, "Metal oxides for optoelectronic applications," *Nature Materials* **15**, 383–396 (2016), 1.
- ²K. Nomura, H. Ohta, A. Takagi, T. Kamiya, M. Hirano, and H. Hosono, "Room-temperature fabrication of transparent flexible thin-film transistors using amorphous oxide semiconductors," *Nature* **432**, 488 (2004).
- ³T. Kamiya, K. Nomura, and H. Hosono, "Origins of High Mobility and Low Operation Voltage of Amorphous Oxide TFTs: Electronic Structure, Electron Transport, Defects and Doping*," *Journal of Display Technology* **5**, 273–288 (2009).
- ⁴T. Kamiya and H. Hosono, "Material characteristics and applications of transparent amorphous oxide semiconductors," *NPG Asia Mater.* **2**, 15 (2010).
- ⁵S. Sallis, K. T. Butler, N. F. Quackenbush, D. S. Williams, M. Junda, D. A. Fischer, J. C. Woicik, N. J. Podraza, B. E. White, A. Walsh, and L. F. J. Piper, "Origin of deep subgap states in amorphous indium gallium zinc oxide: Chemically disordered coordination of oxygen," *Appl. Phys. Lett.* **104**, 232108 (2014).
- ⁶K. M. Niang, J. Cho, S. Heffernan, W. I. Milne, and A. J. Flewitt, "Optimisation of amorphous zinc tin oxide thin film transistors by remote-plasma reactive sputtering," *Journal of Applied Physics* **120** (2016), 10.1063/1.4961608.
- ⁷M. Morales-Masis, F. Dauzou, Q. Jeangros, A. Dabirian, H. Lifka, R. Gierth, M. Ruske, D. Moet, A. Hessler-Wyser, and C. Ballif, "An Indium-Free Anode for Large-Area Flexible OLEDs: Defect-Free Transparent Conductive Zinc Tin Oxide," *Advanced Functional Materials* **26**, 384–392 (2016).
- ⁸E. Nakamura and K. Sato, "Managing the scarcity of chemical elements," *Nature Materials* **10**, 158–161 (2011).
- ⁹J. Heo, S. Bok Kim, and R. G. Gordon, "Atomic layer deposited zinc tin oxide channel for amorphous oxide thin film transistors," *Appl. Phys. Lett.* **101**, 113507 (2012), <http://dx.doi.org/10.1063/1.4752727>.
- ¹⁰W. B. Jackson, R. L. Hoffman, and G. S. Herman, "High-performance flexible zinc tin oxide field-effect transistors," *Appl. Phys. Lett.* **87**, 193503 (2005), <http://dx.doi.org/10.1063/1.2120895>.
- ¹¹S.-M. Lee, Y.-H. Joo, and C.-I. Kim, "Influences of film thickness and annealing temperature on properties of solgel derived ZnOSnO₂ nanocomposite thin film," *Appl. Surf. Sci.* **320**, 494 – 501 (2014).
- ¹²J. H. Ko, I. H. Kim, D. Kim, K. S. Lee, T. S. Lee, B. Cheong, and W. M. Kim, "Transparent and conducting Zn-Sn-O thin films prepared by combinatorial approach," *Applied Surface Science* **253**, 7398–7403 (2007).
- ¹³L.-J. Lee, N.-E. Sung, K. H. Chae, and R. Conley, "Characterization of zinc tin oxide films deposited by radio frequency magnetron sputtering at various substrate temperatures," *Thin Solid Films* **548**, 385 – 388 (2013).
- ¹⁴T. Moriga, Y. Hayashi, K. Kondo, Y. Nishimura, K.-i. Murai, I. Nakabayashi, H. Fukumoto, and K. Tominaga, "Transparent conducting amorphous ZnSnO films deposited by simultaneous dc sputtering," *J. Vac. Sci. Technol., A* **22**, 1705–1710 (2004).
- ¹⁵T. Minami, H. Sonohara, S. Takata, and H. Sato, "Highly Transparent and Conductive Zinc-Stannate Thin Films Prepared by RF Magnetron Sputtering," *Japanese Journal of Applied Physics Part 2-Letters* **33**, L1693–L1696 (1994).
- ¹⁶J. Park, K. T. Oh, D. H. Kim, H. J. Jeong, Y. C. Park, H. S. Kim, and J. S. Park, "High-Performance Zinc Tin Oxide Semiconductor Grown by Atmospheric-Pressure Mist-CVD and the Associated Thin-Film Transistor Properties," *ACS Applied Materials and Interfaces* **9**, 20656–20663 (2017).
- ¹⁷M. K. Jayaraj, K. J. Saji, K. Nomura, T. Kamiya, and H. Hosono, "Optical and electrical properties of amorphous zinc tin oxide thin films examined for thin film transistor application," *Journal of Vacuum Science & Technology B: Microelectronics and Nanometer Structures* **26**, 495 (2008).
- ¹⁸M. Kapilashrami, C. X. Kronawitter, T. Torndahl, J. Lindahl, A. Hultqvist, W.-C. Wang, C.-L. Chang, S. S. Mao, and J. Guo, "Soft X-ray characterization of Zn_{1-x}Sn_xO_y electronic structure for thin film photovoltaics," *Phys. Chem. Chem. Phys.* **14**, 10154–10159 (2012).
- ¹⁹S. C. Siah, S. W. Lee, Y. S. Lee, J. Heo, T. Shibata, C. U. Segre, R. G. Gordon, and T. Buonassisi, "X-ray absorption spectroscopy elucidates the impact of structural disorder on electron mobility in amorphous zinc-tin-oxide thin films," *Appl. Phys. Lett.* **104**, 242113 (2014), <http://dx.doi.org/10.1063/1.4884115>.
- ²⁰J. D. Perkins, J. a. del Cueto, J. L. Alleman, C. Warmsingh, B. M. Keyes, L. M. Gedvilas, P. a. Parilla, B. To, D. W. Readey, and D. S. Ginley, "Combinatorial studies of Zn-Al-O and Zn-Sn-O transparent conducting oxide thin films," *Thin Solid Films* **411**, 152–160 (2002).
- ²¹J. S. Kim, J. K. Park, Y. J. Baik, W. M. Kim, J. Jeong, and T. Y. Seong, "Electrical, optical and etching properties of Zn-Sn-O thin films deposited by combinatorial sputtering," *Journal of the Korean Physical Society* **61**, 1651–1655 (2012).
- ²²P. Görrn, P. Hölzer, T. Riedl, W. Kowalsky, J. Wang, T. Weimann, P. Hinze, and S. Kipp, "Stability of transparent zinc tin oxide transistors under bias stress," *Applied Physics Letters* **90**, 2005–2008 (2007).
- ²³P. Görrn, M. Lehnhardt, T. Riedl, and W. Kowalsky, "The influence of visible light on transparent zinc tin oxide thin film transistors," *Applied Physics Letters* **91**, 2005–2008 (2007).
- ²⁴W. Hu and R. L. Peterson, "Molybdenum as a contact material in zinc tin oxide thin film transistors," *Applied Physics Letters* **104** (2014), 10.1063/1.4875958.

- ²⁵J. Jang, D. G. Kim, D. M. Kim, S.-J. Choi, J.-H. Lim, J.-H. Lee, Y.-S. Kim, B. D. Ahn, and D. H. Kim, "Investigation on the negative bias illumination stress-induced instability of amorphous indium-tin-zinc-oxide thin film transistors," *Applied Physics Letters* **152**108 (2014), 10.1063/1.4898069.
- ²⁶M. G. McDowell, R. J. Sanderson, and I. G. Hill, "Combinatorial study of zinc tin oxide thin-film transistors," *Applied Physics Letters* **92**, 2006–2009 (2008).
- ²⁷A. Marette, A. Poulin, N. Besse, S. Rosset, D. Briand, and H. Shea, "Flexible ZincTin Oxide Thin Film Transistors Operating at 1 kV for Integrated Switching of Dielectric Elastomer Actuators Arrays," *Advanced Materials* **29**, 1–6 (2017).
- ²⁸J. Lee, D.-Y. Cho, J. Jung, U. Ki Kim, S. Ho Rha, C. Seong Hwang, and J.-H. Choi, "Theoretical and experimental studies on the electronic structure of crystalline and amorphous ZnSnO₃ thin films," *Appl. Phys. Lett.* **102**, 242111 (2013).
- ²⁹J. I. Pankove, *Optical Processes in Semiconductors* (Dover Publications, Inc., 1971).
- ³⁰R. Swanepoel, "Determination of the thickness and optical constants of amorphous silicon," *Journal of Physics E: Scientific Instruments* **16**, 1214–1222 (1983).
- ³¹W. L. Bond, "Measurement of the refractive indices of several crystals," *Journal of Applied Physics* **36**, 1674–1677 (1965).
- ³²R. G. Gordon, "Criteria for Choosing Transparent Conductors," *MRS Bulletin* **25**, 5257 (2000).
- ³³E. Arca, K. Fleischer, and I. V. Shvets, "Magnesium, nitrogen codoped Cr₂O₃: A p-type transparent conducting oxide," *Applied Physics Letters* **99**, 111910 (2011).
- ³⁴K. H. L. Zhang, Y. Du, A. Papadogianni, O. Bierwagen, S. Sallis, L. F. J. Piper, M. E. Bowden, V. Shutthanandan, P. V. Sushko, and S. A. Chambers, "Perovskite SrDoped LaCrO₃ as a New pType Transparent Conducting Oxide," *Advanced Materials* **27**, 5191–5195 (2015).
- ³⁵J. H. Jeon, T. K. Gong, Y. M. Kong, H. M. Lee, and D. Kim, "Effect of post-deposition annealing on the structural, optical and electrical properties of IGZO films," *Electronic Materials Letters* **11**, 481–484 (2015).
- ³⁶H. Yanagi, S.-i. Inoue, K. Ueda, H. Kawazoe, H. Hosono, and N. Hamada, "Electronic structure and optoelectronic properties of transparent p-type conducting CuAlO₂," *Journal of Applied Physics* **88**, 4159 (2000).
- ³⁷M. Dekkers, G. Rijnders, and D. H. A. Blank, "ZnIr₂O₄, a p-type transparent oxide semiconductor in the class of spinel zinc-d6-transition metal oxide," *Applied Physics Letters* **90**, 10–13 (2007).
- ³⁸R. Nagarajan, A. D. Draeseke, A. W. Sleight, and J. Tate, "p-type conductivity in CuCr_{1-x}Mg_xO₂ films and powders," *Journal of Applied Physics* **89**, 8022–8025 (2001).
- ³⁹N. Ito, Y. Sato, P. K. Song, A. Kaijio, K. Inoue, and Y. Shigesato, "Electrical and optical properties of amorphous indium zinc oxide films," *Thin Solid Films* **496**, 99–103 (2006).
- ⁴⁰E. Fortunato, D. Ginley, H. Hosono, and D. C. Paine, "Transparent conducting oxides for photovoltaics," *MRS bulletin* **32**, 242–247 (2007).
- ⁴¹A. V. Krukau, O. A. Vydrov, A. F. Izmaylov, and G. E. Scuseria, "Influence of the exchange screening parameter on the performance of screened hybrid functionals," *The Journal of Chemical Physics* **125**, 224106 (2006).
- ⁴²I. Bonalde, E. Medina, M. Rodríguez, S. M. Wasim, G. Marín, C. Rincín, A. Rincín, and C. Torres, "Urbach tail, disorder, and localized modes in ternary semiconductors," *Physical Review B - Condensed Matter and Materials Physics* **69**, 1–5 (2004).
- ⁴³S. John, C. Soukoulis, M. H. Cohen, and E. N. Economou, "Theory of electron band tails and the urbach optical-absorption edge," *Physical Review Letters* **57**, 1777–1780 (1986).
- ⁴⁴M. J. Wahila, K. T. Butler, Z. W. Lebens-Higgins, C. H. Hendon, A. S. Nandur, R. E. Treharne, N. F. Quackenbush, S. Sallis, K. Mason, H. Paik, D. G. Schlom, J. C. Woicik, J. Guo, D. A. Arena, B. E. White, G. W. Watson, A. Walsh, and L. F. J. Piper, "Lone-pair Stabilization in Transparent Amorphous Tin Oxides: A Potential Route to p-type Conduction Pathways," *Chemistry of Materials* **28**, 47064713 (2016).
- ⁴⁵P. J. M. Isherwood, K. T. Butler, A. Walsh, and J. M. Walls, "A tunable amorphous p-type ternary oxide system: The highly mismatched alloy of copper tin oxide," *Journal of Applied Physics* **118**, 105702 (2015).
- ⁴⁶G. Kresse and J. Furthmüller, "Efficient iterative schemes for ab initio total-energy calculations using a plane-wave basis set," *Phys. Rev. B* **54**, 11169–11186 (1996).
- ⁴⁷P. E. Blöchl, "Projector augmented-wave method," *Phys. Rev. B* **50**, 17953–17979 (1994).
- ⁴⁸J. P. Perdew, A. Ruzsinszky, G. I. Csonka, O. A. Vydrov, G. E. Scuseria, L. A. Constantin, X. Zhou, and K. Burke, "Restoring the Density-Gradient Expansion for Exchange in Solids and Surfaces," *Phys. Rev. Lett.* **100**, 136406 (2008).
- ⁴⁹E. S. Goh, J. W. Mah, and T. L. Yoon, "Effects of Hubbard term correction on the structural parameters and electronic properties of wurtzite ZnO," *Computational Materials Science* **138**, 111–116 (2017), arXiv:1703.02496.
- ⁵⁰W. Körner, P. Gumbsch, and C. Elsässer, "Analysis of electronic subgap states in amorphous semiconductor oxides based on the example of Zn-Sn-O systems," *Physical Review B* **86**, 165210 (2012).
- ⁵¹W. Körner and C. Elsässer, "Density-functional theory study of stability and subgap states of crystalline and amorphous Zn-Sn-O," *Thin Solid Films* **555**, 81–86 (2014).
- ⁵²W. Körner, D. F. Urban, and C. Elsässer, "Generic origin of subgap states in transparent amorphous semiconductor oxides illustrated for the cases of In-Zn-O and In-Sn-O," *Physica Status Solidi (a)* **212**, 1476–1481 (2015).
- ⁵³E. Rucavado, Q. Jeangros, D. F. Urban, J. Holovsky, Z. Remes, M. Duchamp, F. Landucci, R. E. Dunin-Borkowski, W. Körner, C. Elsässer, A. Hessler-Wyser, M. Morales-Masis, and C. Ballif, "Enhancing the optoelectronic properties of amorphous zinc tin oxide by subgap defect passivation: A theoretical and experimental demonstration," *Physical Review B* **95**, 1–10 (2017).
- ⁵⁴D. Y. Cho, J. H. Kim, K. D. Na, J. Song, C. S. Hwang, B. G. Park, J. Y. Kim, C. H. Min, and S. J. Oh, "Spectroscopic evidence for limited carrier hopping interaction in amorphous ZnO thin film," *Applied Physics Letters* **95**, 1–4 (2009).
- ⁵⁵D. Schmeißer, J. Haeberle, P. Barquinha, D. Gaspar, L. Pereira, R. Martins, and E. Fortunato, "Electronic structure of amorphous ZnO films," *Physica Status Solidi (C) Current Topics in Solid State Physics* **11**, 1476–1480 (2014).
- ⁵⁶R. Kykyneshi, J. Zeng, and D. P. Cann, "Transparent Conducting Oxides Based on Tin Oxide," in *Handbook of Transparent Conductors*, edited by D. Ginley, H. Hosono, and D. C. Paine (Springer US, 2011) Chap. 6, pp. 171–191.